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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
09/357,507	07/20/1999	KIYOSHI TAGUCHI	10059-286	9338	
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AKIN GUMP STRAUSS HAUER & FELD L.L.P. ONE COMMERCE SQUARE			LEUNG, JENNIFER A		
2005 MARKE	T STREET, SUITE 2200		ART UNIT	ART UNIT PAPER NUMBER	
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Please find below and/or attached an Office communication concerning this application or proceeding.

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	Application No.	Applicant(s)	
Office Action Commence	09/357,507	TAGUCHI ET AL.	
Office Action Summary	Examiner	Art Unit	
	Jennifer A. Leung	1764	
The MAILING DATE of this communication ap Period for Reply	pears on the cover sheet t	with the correspondence address	5
A SHORTENED STATUTORY PERIOD FOR REPL THE MAILING DATE OF THIS COMMUNICATION. - Extensions of time may be available under the provisions of 37 CFR 1. after SIX (6) MONTHS from the mailing date of this communication. - If the period for reply specified above is less than thirty (30) days, a replet If NO period for reply is specified above, the maximum statutory period. - Failure to reply within the set or extended period for reply will, by statut Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	136(a). In no event, however, may a ply within the statutory minimum of the will apply and will expire SIX (6) MC te. cause the application to become the status of the st	i reply be timely filed irty (30) days will be considered timely. NTHS from the mailing date of this commun	ication.
Status			
3) Since this application is in condition for allowa	s action is non-final. ance except for formal ma		its is
closed in accordance with the practice under	Ex parte Quayle, 1935 C.	D. 11, 453 O.G. 213.	
Disposition of Claims			
 4) Claim(s) 1,3,4,6,8-10,21 and 23-28 is/are pen 4a) Of the above claim(s) is/are withdra 5) Claim(s) is/are allowed. 6) Claim(s) 1,3,4,6,8-10,21 and 23-28 is/are rejection is/are objected to. 8) Claim(s) are subject to restriction and/or claim(s) are subject to restriction and/or claim(s) are subject to restriction. 	ected.		
Application Papers			
9) The specification is objected to by the Examine	er.		
10)☐ The drawing(s) filed on is/are: a)☐ acc	cepted or b) objected to	by the Examiner.	
Applicant may not request that any objection to the			
Replacement drawing sheet(s) including the correct 11) The oath or declaration is objected to by the Ex			· ·
Priority under 35 U.S.C. § 119			
12) Acknowledgment is made of a claim for foreign a) All b) Some * c) None of: 1. Certified copies of the priority document 2. Certified copies of the priority document 3. Copies of the certified copies of the priority application from the International Burea * See the attached detailed Office action for a list	ts have been received. ts have been received in a crity documents have been u (PCT Rule 17.2(a)).	Application No received in this National Stage)
Attachment(s)			
Notice of References Cited (PTO-892)	4) Interview	Summary (PTO-413)	
Notice of Draftsperson's Patent Drawing Review (PTO-948) Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08) Paper No(s)/Mail Date	Paper No	s)/Mail Date nformal Patent Application (PTO-152)	

Art Unit: 1764

Response to Amendment

1. Applicant's amendment filed on August 16, 2004 has been received and carefully considered. Claims 2, 5, 7, 11-20 and 22 are cancelled. Claims 1, 3, 4, 6, 8-10, 21 and 23-28 remain active.

Claim Rejections - 35 USC § 102

The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless –

- (b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.
- 2. Claims 1, 6, 8-10, 21 and 25-28 are rejected under 35 U.S.C. 102(b) as being anticipated by James et al. (US 3,262,758).

Regarding claims 1 and 21, James et al. (FIG. 1, 2; generally, column 2, line 24 to column 3, line 60) disclose an apparatus comprising:

- a reaction segment having a catalyst bed for oxidizing carbon monoxide (i.e., the annular space between containers 1 and 8, comprising CO-oxidation bed 22);
- a reformed gas inlet (i.e., conduit 6 for introducing primary reformed gas stream 2);
- a reformed gas pathway for supplying reformed gas to the reaction segment 22 (i.e., indicated by solid-line flow arrows);
- an oxidant gas supplying segment for supplying an oxidant gas to the reformed gas pathway (i.e., conduit 7 for introducing oxygen-containing gas stream 3);
- a cooler for cooling an upstream side of the catalyst bed 22 (i.e., the steam boiler section, located between partitions 18 and 19, comprising conduit 13, fire tubes 17 and heat exchange

Art Unit: 1764

medium supplied via 20 and removed via 21; column 3, lines 25-41); and means for heating a downstream side of the catalyst bed;

wherein the means comprises a portion of the reformed gas pathway located in proximity to and at least partially surrounding the catalyst bed 22; the means being separated from the catalyst bed 22 by a wall (i.e., inner container wall 8), so as to *inherently* heat the downstream side of the bed 22 by the reformed gas and *inherently* cool the reformed gas in the reformed gas pathway via the transfer of heat energy through wall 8 to the catalyst bed 22, before passing through the cooler, via central conduit 13 and fire tubes 17 (column 1, lines 49-71; column 2, lines 57-70; column 3, lines 14-24 and 41-51).

Regarding claims 6 and 25, James et al. further discloses,

"A stream of oxygen containing gas, such as air, is reacted with the primary reformed gas stream. When ammonia synthesis gas is produced, a stoichiometric proportion of air is employed in the secondary reform step to yield a final gas stream containing hydrogen and nitrogen in a 3:1 ratio. The combustion reaction which takes place due to oxygen addition causes a rise in gas stream temperature and some further conversion of hydrocarbon," (column 1, lines 30-38).

Thus, although a gas flow rate control valve is not illustrated in the figures or explicitly disclosed, such control means would be *inherent* of the apparatus, to the enable the disclosed control of a "stoichiometric proportion of air", which will in turn control the "rise in gas stream temperature" and correspond to a rise in the temperature of the catalyst bed 22.

Regarding claims 8 and 26, James et al. (FIG. 1) discloses the reformed gas pathway (see solid-line flow arrows) has a first direction prior to passing through the cooler (i.e. via tube 13) and a second direction passing through the catalyst bed 22, wherein the first and second direction are opposing (column 3, lines 25-52).

Art Unit: 1764

Regarding claim 9, James et al. discloses the reaction segment is located outside the reformed gas pathway (i.e., CO-oxidation bed 22 being annular and having a central reformed gas pathway defined by inner container 8; FIG. 1, 2).

Regarding claim 10, James et al. discloses the reaction segment is tube shaped (i.e., annular, CO-oxidation bed 22; FIG. 1) and the reformed gas pathway before the passage through the cooler (i.e. tubes 13, 17) is formed around the reaction segment 22 (i.e., a portion of the gas stream may be *bypassed around the heat exchange section*, passing out of the lower end of conduit 3 via openings controlled by control dampers 14," column 3, lines 14-24).

Regarding claims 27 and 28, James et al. discloses the portion of the reformed gas pathway (i.e., within inner chamber 8) heats the catalyst bed 22 by direct heat transfer through the wall (column 1, lines 49-71; column 2, lines 58-71).

Instant claims 1, 6, 8-10, 21 and 25-28 structurally read on the apparatus of James et al.

Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103(a), the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 CFR 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103(c) and potential 35 U.S.C. 102(e), (f) or (g) prior art under 35 U.S.C. 103(a).

Art Unit: 1764

3. Claims 3, 4, 23 and 24 are rejected under 35 U.S.C. 103(a) as being unpatentable over James et al. (US 3,262,758) in view of Finneran et al. (US 3,345,136).

James et al. further discloses that in the CO-oxidation stage, "... the cooled gas stream is passed through *one or more beds of promoted iron oxide catalyst*, in order to react carbon monoxide with steam, thus yielding further hydrogen," (column 1, lines 47-49). However, James et al. is silent as to whether the "one or more beds" may be configured such that an upstream side portion of the catalyst is formed of a different catalyst material than the downstream side portion, and whether the catalyst comprising the downstream portion may exert an activity at a lower temperature than the catalyst material of the upstream side portion.

Finneran et al. (column 1, lines 34-61) teaches,

"Regardless of the process for which the hydrogen is made, it is generally desirable that it contain a minimum of residual carbon monoxide," and

"To obtain desirably low levels of residual carbon monoxide in the product gas and a corresponding high degree of conversion to hydrogen, the process can be carried out in two stages. In the first stage, the carbon monoxide containing gas is contacted in the presence of a relatively inexpensive shift conversion catalyst active a relatively high temperatures to convert the bulk of the carbon monoxide. The exothermic heat of the reaction and a substantial portion of the sensible heat of the partially shifted gas is removed by cooling. The cooled gas is then contacted in the presence of a relatively more expensive shift conversion catalyst active a t relatively low temperatures to produce additional hydrogen under equilibrium conditions which favor a low residual proportion of carbon monoxide."

Thus, it would have been obvious for one of ordinary skill in the art at the time the invention was made to select the catalyst configuration as taught by Finneran et al. for the one or more beds of CO-oxidation catalyst 22 in the apparatus of James et al., because the selection of two catalyst stages, wherein an upstream catalyst stage is active at a relatively high temperature and the

Art Unit: 1764

downstream catalyst stage is active at a relatively low temperature, allows a substantial portion of the carbon monoxide to be converted to carbon dioxide, while producing the desirably low levels of residual carbon monoxide in the product gas and a corresponding high degree of conversion to hydrogen, as taught by Finneran et al., above.

Response to Arguments

4. Applicant's arguments submitted on August 16, 2004 have been fully considered but they are not persuasive.

Arguments with respect to JAMES et al.

Beginning on page 3, second to last paragraph, Applicants argue,

"... the reformed gas pathway of James (through chamber 8 and catalyst bed 10) does not "surround" in any way the catalyst bed 22. Instead, it is on the inside of catalyst bed 22, such that catalyst bed 22 surrounds the chamber 8 and catalyst bed 10."

The Examiner maintains her rejection of claim 1, since the reformed gas pathway need only be "located in proximity to said catalyst bed and separated from the catalyst bed by a wall," (lines 8-11). Note that the "surround" feature upon which applicant relies is not recited in the rejected claim 1. Although the claims are interpreted in light of the specification, limitations from the specification are not read into the claims. See *In re Van Geuns*, 988 F.2d 1181, 26

USPQ2d 1057 (Fed. Cir. 1993). Additionally, the Examiner maintains her rejection of claim 21, since the reformed gas pathway need only to "partially surround said catalyst bed," (line 9). As illustrated in FIG. 1, the reformed gas pathway runs from inlet 6 to outlet 25, wherein the pathway partially surrounds catalyst bed 22, for instance, at its upper surface. Note that the feature of a reformed gas pathway that *completely* surrounds the catalyst bed 22 is not recited in the rejected claim 21. Alternatively, one could also argue that the reformed gas pathway does

Art Unit: 1764

completely surround catalyst bed 22, since the reformed gas stream inherently flows through the interstices of catalyst bed 22 within the annular region of the reactor, and therefore "encompasses" the catalyst bed 22.

Beginning on page 4, second paragraph, Applicants argue,

"... James does not intend for the wall of chamber 8 to be a heat transfer wall. Instead, he attempts to avoid such heat transfer by lining the chamber 8 with refractory lining 9."

The Examiner respectfully disagrees. Despite whether or not James intended for heat to transfer through the wall of chamber 8, the Examiner maintains that the refractory lined wall of chamber 8 will inherently heat the downstream side of the catalyst bed 22. James specifically addresses this assertion by disclosing,

"... the bed of CO-oxidation catalyst acts as a temperature moderator and serves as insulation to protect the outer container from the high temperature maintained in the secondary converter section." (column 1, lines 57-61; also, column 2, lines 13-16).

Heat is necessarily transferred through the refractory lining 9 and the wall of chamber 8, in order for the bed of CO-oxidation catalyst 22 to exhibit its disclosed temperature moderating and insulating characteristics.

The generally known nature of heat transfer in refractory materials is evidenced by White. White discloses an apparatus for heating a heat transfer fluid, such as water, wherein the apparatus comprises a wall including a refractory layer 9 that separates a higher-temperature firebox 4 from a lower-temperature water jacket 13 (FIG. 2). In particular, White (page 7, line 17, to page 8, line 8) teaches that,

"... due to the nature of refractory layer 9 significant heat build-up may be achieved within the layer 9 over prolonged periods whereby layer 9 then functions as a heat reservoir wherein heat may be transferred to the water contained within the water jacket

Art Unit: 1764

13 adjacent the layer 9 thereby providing a secondary region of heat transfer, i.e. a preheating area."

One having ordinary skill in the art would recognize that the refractory lining 9 in the apparatus of James would exhibit heat transfer characteristics similar to the refractory lining 9 in the apparatus of White. In the apparatus of James, significant heat build-up would similarly occur within the refractory layer 9 over prolonged periods, whereby layer 9 would then function as a heat reservoir wherein heat may be transferred to the adjacent stainless steel wall of chamber 8 and subsequently into the adjacent layer of catalyst 22, thereby creating a heating means for the downstream side of catalyst 22 and causing the catalyst 22 to exhibit its temperature moderating and insulating characteristics.

Beginning on page 4, third paragraph, Applicants argue,

"... it cannot be said that the temperature of the catalyst bed 22 necessarily increases from the upstream side to the downstream side, i.e., is heated at the downstream side... there are some cases where the temperature increases from the downstream side to the upstream side of the catalyst bed 10. In such a case, the catalyst bed 22 could be heated at the upstream end and cooled by heat lost from the catalyst bed 10. Therefore, the Examiner's contention of inherency is again incorrect, because heating and cooling alleged by the Examiner do not necessarily occur in the James apparatus."

The Examiner respectfully disagrees. As disclosed by James (column 2, lines 58-70),

"The mixed process gases react in the combustion chamber. The resulting combustion will raise the process gas stream temperature from 1300° F. to about 2000° F. The hot gas stream now rises from chamber 4, passing to inner container 8 provided with refractory lining 9... Bed 10 will typically be at a temperature of about 1650° F., and final catalytic conversion of unreacted hydrocarbon in the gas stream thus is accomplished in bed 10."

Looking now to FIG. 1, the first region located immediately upstream of bed 10 within chamber 8 thereby contains process gas at about 2000° F. The second region within bed 10 between

Art Unit: 1764

retention means 11 and 12 thereby contains process gas undergoing reaction, at about 1650° F. And the third region located immediately downstream of bed 10 within chamber 8 can be assumed to contain process gas at a temperature no greater than 1650° F, since no catalyst is provided within this region. One of ordinary skill in the art would observe that the temperature profile from the first region (~2000° F) to the second region (~1650° F) to the third region (≤1650° F) within chamber 8 progressively decreases. Subsequently, the CO-oxidation catalyst 22 that is located adjacent to chamber 8 will inherently be heated by a corresponding region of process gas, with the temperature of catalyst bed 22 increasing from its upstream side (i.e., adjacent to the third region of about ≤1650° F, and the 700° F inlet stream; column 3, lines 42-44) to its downstream side (i.e., adjacent to the first region of about 2000° F).

Arguments with respect to JAMES et al. in view of FINNERAN, JR. et al.

Beginning on page 7, second paragraph, Applicants argue,

"Applicants do not contest that the conversion of residual carbon monoxide in two stages with catalysts active at different temperatures is generally known. However, the Examiner does not suggest how the teachings of Finneran would be incorporated in the apparatus of James."

In response to applicant's argument that there is no suggestion to combine the references, the examiner recognizes that obviousness can only be established by combining or modifying the teachings of the prior art to produce the claimed invention where there is some teaching, suggestion, or motivation to do so found either in the references themselves or in the knowledge generally available to one of ordinary skill in the art. See *In re Fine*, 837 F.2d 1071, 5 USPQ2d 1596 (Fed. Cir. 1988) and *In re Jones*, 958 F.2d 347, 21 USPQ2d 1941 (Fed. Cir. 1992). Additionally, the test for obviousness is not whether the features of a secondary reference

Art Unit: 1764

Page 10

may be bodily incorporated into the structure of the primary reference; nor is it that the claimed invention must be expressly suggested in any one or all of the references. Rather, the test is what the combined teachings of the references would have suggested to those of ordinary skill in the art. See *In re Keller*, 642 F.2d 413, 208 USPQ 871 (CCPA 1981). The teachings of Finneran Jr., et al. would have suggested to one of ordinary skill in the art that modifying the one or more catalyst bed 22 in the apparatus of James et al. to comprise an upstream side portion of different material and activity than the downstream side portion would have allowed a substantial portion of the carbon monoxide to be converted to carbon dioxide, while producing the desirably low levels of residual carbon monoxide in the product gas and a corresponding high degree of conversion to hydrogen, in addition to minimizing the expense of catalyst. (see rejection above).

Conclusion

5. THIS ACTION IS MADE FINAL. Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of this final action.

Art Unit: 1764

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Jennifer A. Leung whose telephone number is (571) 272-1449. The examiner can normally be reached on 8:30 am - 5:30 pm M-F, every other Friday off.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Glenn A. Caldarola can be reached on (571) 272-1444. The fax phone number for the organization where this application or proceeding is assigned is 703-872-9306.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

Jennifer A. Leung November 4, 2004

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PRINARIMARY EXAMINER